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THE CLAIMS DEFINING THE INVENTION ARE AS FOLLOWS:

- 1. A method of treating a drug resistant disease in a subject in need thereof comprising the step of administering to said subject a therapeutically effective amount of hyaluronan in conjunction with a chemotherapeutic agent such that said chemotherapeutic agent is more effective than when administered alone.
- 2. A method of enhancing the bioavailability of a chemotherapeutic agent comprising the step of administering to a subject in need thereof a therapeutically effective amount of hyaluronan.
- 3. A method of treating or preventing multidrug resistance or drug-resistant cells comprising the step of administering a sufficient amount of hyaluronan, prior to, together with, or subsequent to the administration of a chemotherapeutic jagent.
- 4. A method according to any one of claims 1 to 3, wherein the chemotherapeutic agent is selected from the 20 group consisting of carmustine (BCNU), chlorambucil (Leukeran), cisplatin (Platinol), Cytarabine, doxorubicin (Adriamycin), fluorouracil (5-FU), methoxetrate (Mexate), CPT111, etoposide, plicamycin (Mithracin) and taxanes.
- 5. A method according to claim 1, wherein the drug resistant disease is a cellular proliferative disorder.
 - 6. A method according to claim 5, wherein the cellular proliferative disorder is selected from the group consisting of cancers of the breast, lung, prostate, kidney, skin, neural, ovary, uterus, liver, pancreas,
- 30 epithelial, gastric, intestinal, exocrine, endocrine, lymphatic, hematopoietic system or head and neck tissue.
 - 7. A method according to any one of claims 1 to 6, wherein the subject is mammal.
- 8. A method according to claim 7, wherein the mammal is selected from the group consisting of bovine, canine, equine, feline, porcine and human.
 - 9. A method according to any one of claims 1 to 8,

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wherein the administration of hyaluronan, is prior to, together with, or subsequent to the administration of a chemotherapeutic agent.

- 10. A method according to any one of claims 1 to 9, 5 wherein the administration of hyaluronan and/or chemotherapeutic agent is orally, topically, or parenterally.
 - 11. A method according to claim 10, wherein the hyaluronan and or chemotherapeutic agent is administered together with a pharmaceutically acceptable carrier, adjuvant, or vehicle.
 - 12. A method according to claim 10 or claim 11, wherein parenteral administration is either by subcutaneous injection, aerosol, intravenous, intramuscular,
- intrathecal, intracranial, intrasternal injection or infusion techniques.

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